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NEWS 8 JAN 17 IPC 8 in the WPI family of databases including WPIFV
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NEWS 12 FEB 22 Status of current WO (PCT) information on STN
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NEWS 14 FEB 22 Updates in EPFULL; IPC 8 enhancements added
NEWS 15 FEB 27 New STN AnaVist pricing effective March 1, 2006
NEWS 16 FEB 28 MEDLINE/LMEDLINE reload improves functionality
NEWS 17 FEB 28 TOXCENTER reloaded with enhancements
NEWS 18 FEB 28 REGISTRY/ZREGISTRY enhanced with more experimental spectral
property data
NEWS 19 MAR 01 INSPEC reloaded and enhanced
NEWS 20 MAR 03 Updates in PATDPA; addition of IPC 8 data without attributes
NEWS 21 MAR 08 X.25 communication option no longer available after June 2006
NEWS 22 MAR 22 EMBASE is now updated on a daily basis

NEWS EXPRESS FEBRUARY 15 CURRENT VERSION FOR WINDOWS IS V8.01a,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
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 75316 SULFONIC
 (SULFONIC OR SULFONICS)
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1516900 ACIDS
4612947 ACID
 (ACID OR ACIDS)
72735 SULFONIC(L)ACID
L2 2 L1 AND (SULFONIC(L)ACID)

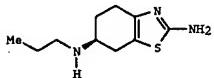
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L2 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2006:31539 CAPLUS
 DOCUMENT NUMBER: 144:128965
 TITLE: Improved process for the preparation of biologically active tetrahydrobenzothiazole derivative
 INVENTOR(S): Mistry, Dhiren N.; Soni, Kamlesh S.; Vasoya, Sanjay L.; Kansal, Vinod Kumar
 PATENT ASSIGNEE(S): Alembic Limited, India
 SOURCE: PCT Int. Appl., 31 pp.
 CODEN: PIXDD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|----------|
| WO 2006003677 | A1 | 20060112 | WO 2005-IN127 | 20050425 |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BE, BG, BR, BY, CZ, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LX, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TH, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW | | | | |
| RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BE, BY, CF, CG, CI, CM, GA, GN, GQ, GW, HL, MR, NE, SN, TD, TG, BV, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZV, AM, AZ, BY, KG, KZ, HD, RU, TJ, TM | | | | |
| PRIORITY APPLN. INFO.: IN 2004-MU706 | A | 20040701 | | |

GI



I

AB A process for an improved preparation of Pramipexole (I) is described. The process begins with 4-aminocyclohexanol which is protected with a phthalimido group then subjected to oxidation, bromination, cyclocondensation with thiourea, deprotected, resolved with tartaric acid, and finally reductive alkylation with propionaldehyde.
 REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2000:725436 CAPLUS
 DOCUMENT NUMBER: 133:301171
 TITLE: Compositions and methods for improved delivery of ionizable hydrophobic therapeutic agents
 INVENTOR(S): Chen, Feng-jing; Patel, Manesh V.
 PATENT ASSIGNEE(S): Lipocene, Inc., USA
 SOURCE: PCT Int. Appl., 99 pp.
 CODEN: PIXKD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|----------|
| WO 2000059475 | A1 | 20001012 | WO 2000-US7342 | 20000316 |
| W: AE, AL, AM, AT, AU, AZ, BA, BE, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KW, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM | | | | |
| RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IR, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CN, GA, GN, GQ, GW, HL, MR, NE, SN, TD, TG, BV, GH, GM, Tween-20 | | | | |
| US 6383471 | BI | 20020507 | US 1999-287043 | 19990406 |
| CA 2366702 | AA | 20001012 | CA 2000-2366702 | 20000316 |
| EP 1165048 | A1 | 20020102 | EP 2000-916547 | 20000316 |
| PRIORITY APPLN. INFO.: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO | | | | |

US 1999-287043 A 19990406
 WO 2000-US7342 W 20000316

AB The present invention is directed to a pharmaceutical composition including a

hydrophobic therapeutic agent having at least one ionizable functional group, and a carrier. The carrier includes an ionizing agent capable of ionizing the functional group, a surfactant, and optionally solubilizers, triglycerides, and neutralizing agents. The invention further relates to a method of preparing such compns. by providing a composition of an ionizable

hydrophobic therapeutic agent, an ionizing agent, and a surfactant, and neutralizing a portion of the ionizing agent with a neutralizing agent. The compns. of the invention are particularly suitable for use in oral dosage forms. A carrier containing concentrated phosphoric acid 0.025, Tween-20 0.3, Ariacel 186 0.2, sodium taurocholate 0.15, propylene glycol 0.3 g was formulated. Itraconazole was included in the carrier at 30 mg/mL for testing the stability of the itraconazole solution upon dilution in simulated gastric fluid.

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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